

DG DEEP CLEANING ASTRINGENT SENSITIVE SKIN- salicylic acid liquid
BELVEDERE INTERNATIONAL INC.

Reference Label Set Id: 711e7953-33f1-48ad-af74-16f25341c404

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

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Active Ingredient

Salicylic acid 0.50%

USE for treatment of acne

Keep out of reach of children, if swallowed, get medical help or contact a Poison Control Center immediately.

USE FOR TREATMENT OF ACNE

FOR EXTERNAL USE ONLY, Flammable, keep away from open fire or flame, Using other topical acne medication at the same time or immediately following the use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor. Rinse right away with water if it gets in eyes.

- CLEANSE SKIN THOROUGHLY BEFORE APPLYING MEDICATION
- MOISTEN A COTTON BALL AND COVER THE ENTIRE AFFECTED AREA WITH A THIN LAYER 1 TO 3 TIMES DAILY
- BECAUSE EXCESSIVE DRYING OF THE SKIN MAY OCCUR, START WITH 1 APPLICATION DAILY. THEN GRADUALLY INCREASE TO 2 OR 3 TIMES DAILY IF NEEDED OR AS DIRECTED BY A DOCTOR. IF BOTHERSOME DRYING OR PEELING OCCURS, REDUCE APPLICATION TO ONCE A DAY OR EVERY OTHER DAY.

WATER, Alcohol, Glycerin, Isoceteth-20, Sodium Citrate, Fragrance, Propylene Glycol, Benzophenone-4, Algae extract, Aloe barbadensis leaf extract, triethanolamine, denatonium benzoate, blue 1



Deep Cleaning Astringent Sensitive Skin

Salicylic Acid
Acne Medication

Compare to Clean & Clear®
Deep Cleaning Astringent*

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Purpose

Acne medication

Use for treatment of acne

Warnings

For external use only

Flammable, keep away from open fire or flame

■ Using other topical acne medication at the same time or immediately following the use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor

■ Rinse right away with water if it gets in eyes.

Keep out of reach of children, if swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- cleanse skin thoroughly before applying medication
- moisten a cotton ball and cover the entire affected area with a thin layer 1 to 3 times daily
- because excessive drying of the skin may occur, start with 1 application daily, then gradually increase to 2 or 3 times daily if needed or as directed by a doctor. If bothersome drying or peeling occurs, reduce application to once a day or every other day.

Inactive ingredients water, alcohol, glycerin, isoceteth-20, sodium citrate, fragrance, propylene glycol, benzophenone-4, algae extract, aloe barbadensis leaf extract, triethanolamine, denatonium benzoate, blue 1

Other information Store at room temperature

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Clean & Clear®.

**100%
Quality
Guaranteed**
(888) 309-9030

B0106



DISTRIBUTED BY
DOLGENCORP, LLC
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

MADE IN CANADA LB80001B

DG DEEP CLEANING ASTRINGENT SENSITIVE SKIN

salicylic acid liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:60742-002

Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.5 g in 100 g	
Inactive Ingredients				
	Ingredient Name	Strength		
	WATER (UNII: 059QF0KO0R)			
	ALCOHOL (UNII: 3K9958V90M)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	ISO CETETH-20 (UNII: O020065R7Z)			
	SODIUM CITRATE (UNII: 1Q73Q2JULR)			
	TROLAMINE (UNII: 9O3K93S3TK)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	BENZOPHENONE (UNII: 701M4TTV9O)			
	ALOE (UNII: V5VD430YW9)			
	DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)			
	FD&C BLUE NO. 1 (UNII: HBR47K3TBD)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60742-002-01	240 g in 1 BOTTLE		
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph not final	part333D	01/31/2013	

Labeler - BELVEDERE INTERNATIONAL INC. (247836356)

Registrant - BELVEDERE INTERNATIONAL INC. (247836356)

Establishment

Name	Address	ID/FEI	Business Operations
BELVEDERE INTERNATIONAL INC.		247836356	manufacture(60742-002)